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Case 3:07-cv-06352-MMC

PLEASE TAKE NOTICE that defendant Medtronic, Inc. ("Medtronic") hereby removes the above-captioned action from the Superior Court of the State of California in and for the County of San Francisco to the United States District Court for the Northern District of California pursuant to 28 U.S.C. §§ 1332 and 1441. As set forth more fully below, this case is properly removed to this Court pursuant to 28 U.S.C. § 1441 because Medtronic has satisfied the procedural requirements for removal and this Court has subject matter jurisdiction over this action pursuant to 28 U.S.C. § 1332. In support of this Notice of Removal, Medtronic states the following:

#### I. THE PROCEDURAL REQUIREMENTS FOR REMOVAL ARE SATISFIED

- 1. On November 5, 2007, Plaintiffs Roseann Mitchell, Uriela Mitchell, Thomas Mitchell and Patricia Hines ("Plaintiffs") commenced this action entitled *Mitchell, et al. v. Medtronic, Inc., et al.*, Case No. CGC-07-468894 (the/this "Action") against Medtronic and McKesson Corporation ("McKesson") by filing a Complaint (the "Complaint") with the Superior Court of the State of California in and for the County of San Francisco. Plaintiffs' Complaint purports to plead a number of causes of action, but all of Plaintiffs' putative claims are premised on the basic allegation that Roseann Mitchell was injured by allegedly defective Medtronic Sprint Fidelis cardiac-defibrillator leads ("Fidelis Leads"). A true and correct copy of the Complaint and Summons are attached hereto as Exhibit "A".
- 2. The Complaint was served on Medtronic on November 15, 2007. Medtronic is informed and believes that defendant McKesson was served on November 15, 2007. Accordingly, this Notice of Removal is timely filed pursuant to 28 U.S.C. § 1441.
  - 3. All properly joined and served defendants consent to this removal.<sup>1</sup>

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<sup>1</sup> The Complaint purports to name McKesson as codefendant. As discussed below, however, McKesson has been fraudulently joined as a defendant herein. Accordingly, McKesson's consent to this removal is not required. See 28

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The Superior Court of the State of California in and for the County of San Francisco 4. is located within the Northern District of California. See 28 U.S.C. § 83. Thus, venue is proper in this Court because it is the "district and division embracing the place where such action is pending." 28 U.S.C. § 1441(a).

- On December 13, 2007 Medtronic filed an Answer to the Complaint. A true and correct copy of Medtronic's Answer filed in this action is attached hereto as Exhibit "B."
  - No previous application has been made for the relief requested herein. 6.
- Pursuant to 28 U.S.C. § 1446(a), a copy of all process, pleadings, and orders served 7. upon Medtronic, including the Complaint and Summons, are attached hereto.
- Pursuant to 28 U.S.C. § 1446(d), a copy of this Notice of Removal is being served 8. upon counsel for Plaintiffs and a copy is being filed with the Clerk of the Court for the Superior Court of the State of California in and for the County of San Francisco.2
- REMOVAL IS PROPER BECAUSE THIS COURT HAS SUBJECT MATTER II. **JURISDICTION PURSUANT TO 28 U.S.C. § 1332**
- This Court has subject matter jurisdiction pursuant to 28 U.S.C. § 1332 because this 9. action is between citizens of different states and it is apparent that Plaintiffs seek recovery of an amount in excess of \$75,000, exclusive of costs and interest.

U.S.C. § 1441(b); Hewitt v. City of Stanton, 798 F.2d 1230, 1233 (9th Cir. 1986) (co-defendants who are fraudulently joined need not join in removal).

<sup>2</sup> By filing a Notice of Removal in this matter, Medtronic does not waive its right to object to service of process, the sufficiency of process, jurisdiction over the person, or venue, and Medtronic specifically reserves the right to assert any defenses and/or objections to which it may be entitled.

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#### A. **Diversity of Citizenship**

- 11. Medtronic is now, and at the time of the filing of this action was, a corporation existing under the laws of the State of Minnesota, having its principal place of business in the State of Minnesota. See Complaint at ¶ 6 (alleging that Medtronic "is a Minnesota corporation" with its principal place of business in Minneapolis, Minnesota). Medtronic thus is a citizen of Minnesota for purposes of determining diversity. 28 U.S.C. § 1332(c)(1).
- 12. Plaintiffs' Complaint alleges that Plaintiffs are citizens of the State of California. (Complaint at ¶¶2-5). Based on this allegation, Medtronic is informed and believes that Plaintiffs are, and at the time of the filing of this action were, citizens of the State of California. 30 M.
- 13. Upon information and belief, none of the Doe defendants have been substituted with any named defendants or been served with process in the state court action. For purposes of removal, "the citizenship of defendants sued under fictitious names shall be disregarded." 28 U.S.C. §1441(a); accord Soliman v. Phillip Morris Inc., 311 F.3d 966, 971 (9th Cir. 2002); McCabe v. General Foods Corp., 811 F.2d 1336, 1339 (9th Cir. 1987). Therefore, the citizenship of Does 1 to 50 should be disregarded for purposes of diversity.
- 14. Because the remaining named defendant – McKesson – is fraudulently joined, its citizenship must likewise be ignored for the purpose of determining the propriety of removal. See McCabe, 811 F.2d at 1339. Accord Ritchey v. Upjohn Drug Co., 139 F.3d 1313, 1318 (9th Cir. 1998) ("It is a commonplace that fraudulently joined defendants will not defeat removal on diversity grounds.") (citations omitted). 11

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- As the Ninth Circuit has explained, "[f]raudulent joinder is a term of art." McCabe, 15. 811 F.2d at 1339.3 A defendant is fraudulently joined and its presence in the lawsuit is ignored for purposes of determining diversity where no viable cause of action has been stated against the resident defendant. See Morris v. Princess Cruises, Inc., 236 F.3d 1061, 1067 (9th Cir. 2001); Ritchey, 139 F.3d at 1318-19; United Computer Sys., Inc. v. AT&T Corp., 298 F.3d 756, 761 (9th Cir. 2002).
- A defendant is fraudulently joined and its presence in the lawsuit is ignored for 16. purposes of determining diversity "if the plaintiff fails to state a cause of action against the resident defendant, and the failure is obvious according to the settled rules of the state." Morris, 236 F.3d at 1067; Ritchey, 139 F.3d at 1318-19; TPS Utilicom Services Inc., 223 F. Supp. 2d 1089, 1100 (C.D. Cal. 2002) (citations omitted).
- Here, it is obvious on the face of the Complaint that defendant McKesson has been 17. fraudulently joined because Plaintiffs do not make any material factual allegations against McKesson at all. The Supreme Court has made clear that when filing a complaint, "a plaintiff's obligation to provide the grounds of his entitlement to relief requires more than labels and conclusions, and a formulaic recitation of the elements of a cause of action will not do." Bell Atlantic v. Twombly, 127 S.Ct. 1955, 1964-65 (2007). The plaintiff must provide some factual allegations of the grounds upon which the claim rests (id. at 1965 n. 3) and the failure to assert material factual allegations against a defendant is a hallmark of fraudulent joinder. See, e.g., Taylor v. Shetler Lincoln Mercury Ltd., 2007 WL 3244701, No. 2:07-CV-0097, \* 1-2 (W.D. La. Nov. 2, 2007) (citing Twombly and finding fraudulent joinder); Pascale Service Corp. v. International Truck And Engine Corporation, 2007 WL 2905622, No. 07-0247-S, \*2-4 (D.R.I. Oct. 1, 2007) (citing Twombly and finding fraudulent joinder); Lyons v. American Tobacco Co., No. Civ. A. 96-0881-BH-

It "is not intended to impugn the integrity of Plaintiff[s] or [their] counsel." Brown v. Allstate Ins. Co., 17 F. Supp.2d 1134, 1137 (S.D. Cal. 1998).

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S, 1997 WL 809677, at \*5 (S.D. Ala. Sept. 30, 1997) (holding that there is "no better admission of fraudulent joinder of [the resident defendants]" than the failure of the plaintiff "to set forth any specific factual allegations" against them).

18. All that Plaintiffs offer here as to McKesson is a single boilerplate allegation that, as a general matter, McKesson markets, distributes, promotes, advertises, and sells Fidelis Leads. (Complaint at ¶7). But just as in Lyons, there are "no specific factual allegations in the Complaint to support such claims against the distributor[] [defendant]." 1997 WL 809677, at \*5. Nor do Plaintiffs allege any facts that would indicate that McKesson sold or distributed the Fidelis Leads that were allegedly implanted in the Plaintiff.<sup>4</sup> Plaintiffs cannot cure these deficiencies by simply relying on allegations directed toward "defendants." See Lyons, 1997 WL 809677, at \*5 ("Although plaintiffs in their complaint as in their respective briefs filed in support of their motion to remand. generally refer to all defendants collectively with respect to at least three of their causes of action. there exists no specific factual allegations in the Complaint to support such claims against the distributors."); In re Phenylpropanolamine (PPA) Prod. Liab. Litig., ("In re PPA"), MDL No. 1407, Docket No. C02-423R, Slip. Op. at 5 (W.D. Wash. Nov. 27, 2002) (a true and correct copy is attached hereto as Exhibit "C") (allegations directed toward "defendants" or "all defendants" insufficient). Accordingly, on its face, the Complaint fails to state a viable cause of action against McKesson.

19. Moreover, Plaintiffs' failure to allege any facts to support a claim against McKesson is not a mere pleading deficiency - Plaintiffs cannot allege that McKesson sold or distributed the Sprint Fidelis leads allegedly at issue in this case.

<sup>4</sup> Indeed, although the crux of Plaintiffs' Complaint is an alleged failure to adequately warn of alleged risks associated with the Fidelis leads, the Complaint does not include any allegations that McKesson made any specific representations or warranties to plaintiff's implanting physician, or that plaintiff or plaintiff's physicians relied on any such specific representations or warranties by McKesson.

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- 20. The Ninth Circuit has made clear that "[w]here fraudulent joinder is an issue, . . . '[t]he defendant seeking removal to the federal court is entitled to present the facts showing the joinder to be fraudulent." Ritchey, 139 F.3d at 1318 (quoting McCabe, 811 F.2d at 1339). Thus, "If raudulent joinder claims may be resolved by 'piercing the pleadings' and considering summary judgment-type evidence such as affidavits and deposition testimony." Morris, 236 F.3d at 1068 (citation omitted). Accord Ritchey, 139 F.3d at 1318 ("If we had been required [in McCabe] to look at facts outside of the complaint to decide that issue, we would have done so."); McCabe, 811 F.2d at 1339 (considering sworn Declarations in determining fraudulent joinder issue).
- 21. Thus, "a removing defendant" may "submit facts showing that a resident defendant had 'no real connection with the controversy." Ritchey, 139 F.3d at 1318 (quoting Wilson v. Republic Iron & Steel Co., 257 U.S. 92, 97 (1921)). That is precisely the situation here. McKesson is not properly joined as a defendant herein because as the Declaration submitted by Medtronic establishes, McKesson is not now, and has never been, an authorized seller or distributor of Sprint Fidelis Leads in the State of California or anywhere else in the United States and, as such, McKesson did not distribute or sell any Sprint Fidelis lead at issue in this action. (See Affidavit of Henry David Roberts ("Roberts Aff.), attached hereto as Exhibit "D"). Indeed, McKesson has never been authorized to market, distribute, promote, advertise, or sell Medtronic Sprint Fidelis Leads in the State of California or anywhere else in the United States. Id.
- 22. Accordingly, McKesson was improperly joined as a defendant and its citizenship may not be considered in assessing whether there is diversity of citizenship under 28 U.S.C. § 1332.
- 23. Because Plaintiffs allegedly are citizens of California and the sole properly named Defendant, Medtronic, is not a citizen of California, complete diversity exists in this action.

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whether it is 'facially apparent' that the claims exceed the jurisdictional amount." White v. FCI USA, Inc., 319 F.3d 672, 675 (5th Cir. 2003).

25. Plaintiffs here allege injuries as a result of Roseann Mitchell's implantation with Fidelis leads (Complaint at ¶¶ 14, 18, 21, 27, 83). Plaintiffs do not seek to limit the amount of compensatory damages that they allege they are entitled to receive.

26. Plaintiffs allege that the Sprint Fidelis leads were defective, that as a result "she experienced multiple episodes of severe and obvious shock to her heart and person," and that "[a]s a consequence therein, she underwent explant surgery at which the physicians discovered that the Sprint Fidelis Lead, model 6949, had fractured and the shocks were a result of said fracture." (Complaint, ¶ 14.) See also id., ¶ 21 (alleging that Plaintiff Roseann Mitchell was "forced to have an early explant and implant of a new lead system, scarring her already fragile heart, and forcing her to undergo additional and unnecessary complicated surgery" and that "[a]s a result," she "has suffered severe physical and emotional injuries . . . in addition to the injuries and damages sustained as a result of the shocks that preceded the explantation"); id, ¶ 27 (alleging that "[a]s a result of the fracture" of the Sprint Fidelis leads, Plaintiff Roseann Mitchell "suffered inappropriate shocking leading to the need for dangerous and serious cardiac surgery, required and will continue to require healthcare and services; has incurred and will continue to incur medical and related expenses; has suffered and will continue to suffer mental anguish, diminished capacity for the enjoyment of life, a diminished quality of life, increased risk of premature death, and other such damages"). And, finally, Plaintiffs allege that their damages include, inter alia, general damages, damages for

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medical, hospital, and incidental expenses, loss of earnings and loss of earning capacity, and punitive and exemplary damages. (Complaint at ¶ 85).

- 27. Based on these allegations, it is "facially apparent" from their Complaint that Plaintiffs seek an amount in excess of \$75,000.
- 28. Other federal courts have found that the amount in controversy likely exceeds \$75,000 in cases alleging similar types of damages. See, e.g., Luckett v. Delta Airlines, Inc., 171 F.3d 295, 298 (5th Cir. 1999) (finding it "facially apparent" that plaintiff's claim exceeded \$75,000 where plaintiff alleged property damage, travel expenses, an emergency ambulance trip, a six-day stay in the hospital, pain and suffering, humiliation, and temporary inability to do housework after her hospitalization); White, 319 F.3d at 674 (finding it "facially apparent" that plaintiff's wrongful termination claim exceeded \$75,000 based on claims for loss of pay, fringe benefits, impaired earning capacity, harm to credit, emotional distress, attorney fees, punitive damages, etc.). The District Court in White concluded that the compensatory damages alone "in all likelihood" exceeded \$75,000. White, 319 F.3d at 675.

WHEREFORE, Medtronic respectfully removes the above-described action from the Superior Court of the State of California in and for the County of San Francisco to this Court.

'DATED: December 14, 2007.

REED SMITH LLP

Attorneys for Defendant

Medtronic Inc.

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#### **DEMAND FOR JURY TRIAL**

1.

Plaintiffs herewith request a trial by jury as to all issues of material fact.

#### **PARTIES**

2.

Plaintiff ROSEANN MITCHELL is, and was, at all relevant times, a resident of the State of California.

3.

Plaintiff URIELA MITCHELL is, and was, at all relevant times, a resident of the State of California and is the daughter of Plaintiff ROSEANN MITCHELL.

4.

Plaintiff PATRICIA HINES is, and was, at all relevant times, a resident of the State of California and is the daughter of Plaintiff ROSEANN MITCHELL.

5.

Plaintiff THOMAS MITCHELL is, and was, at all relevant times, a resident of the State of California and is the son of Plaintiff ROSEANN MITCHELL.

6.

Defendant MEDTRONIC, INC. is a Minnesota corporation, with its principal place of business at 710 Medtronic Parkway, Minneapolis, Minnesota 55432. Medtronic develops technology to treat conditions such as heart disease and other illnesses. Medtronic manufactures, markets, advertises, promotes and sells Medtronic implantable defibrillators ("ICDs"), and "SPRINT FIDELIS Leads," worldwide.

7.

Defendant McKESSON CORPORATION, ("McKESSON") is a corporation existing under the laws of incorporation of the State of New Jersey, with its principal

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place of business in San Francisco, California. At all times herein mentioned, Defendant McKESSON was, and is, been engaged in the business of marketing, distributing, promoting, advertising and selling MEDTRONIC SPRINT FIDELIS Leads nationwide and in the State of California.

8.

In engaging in the conduct alleged herein, each Defendant acted as the agent for each of the other Defendants.

9.

Defendants MEDTRONIC, INC., McKESSON and Does One through Fifty will hereafter be referred to as "Defendants".

10.

At all times relevant to this action, Defendants intentionally, recklessly and/or negligently concealed, suppressed, omitted, and misrepresented the risks, dangers, defects, and disadvantages of Medtronic SPRINT FIDELIS Leads, and advertised, promoted, marketed, sold and distributed Medtronic SPRINT FIDELIS Leads as a safe medical device when, in fact, Defendants knew that Medtronic SPRINT FIDELIS Leads were not safe for their intended purposes and that Medtronic SPRINT FIDELIS Leads would cause, and did cause, serious medical problems, and in some patients, catastrophic injuries and deaths.

11.

At all relevant times herein, Defendants, at all times relevant herein, designed, developed, manufactured, promoted, marketed, distributed, tested, warranted and sold in interstate commerce (including California) SPRINT FIDELIS Leads. Defendant McKesson has its principal place of business in San Francisco, California, and all said defendants do substantial business in the State of California, advertise in California, and receive substantial compensation and profits from sales of Medtronic SPRINT FIDELIS Leads in

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California.

#### FACTUAL ALLEGATIONS

12.

Defendants design, research, develop, manufacture, test, market, advertise, promote, distribute, and sell products that are sold to and marketed to treat cardiac arrhythmias, heart failure, and coronary and peripheral vascular disease including implantable cardiac defibrillators ("ICDs") and the SPRINT FIDELIS Leads which are the subject of this lawsuit. ICD detects and corrects both fast and slow heart rates. The defibrillator portion regulates the heart through electrical signals and allows an appropriate rhythm to take over. The SPRINT FIDELIS Leads connect the ICD to the heart through a blood vessel and are intended to transmit electrical impulses that are necessary to maintain normal rate and rhythm.

13.

Plaintiff ROSEANN MITCHELL has a cardiovascular condition that necessitates the use of an ICD and was implanted with such device on December 31, 2005. The ICD was attached to her heart with the lead wire system called SPRINT FIDELIS Lead, model number 6949, designed, developed, manufactured, marketed, promoted, advertised, tested, warranted and distributed by defendants herein.

14.

Between early 2006 through September 2007, Plaintiff ROSEANN MITCHELL experienced episodes of shock and in September 2007, while in the car with the other plaintiffs herein, she experienced multiple episodes of severe and obvious shock to her heart and person. As a consequence therein, she underwent explant surgery at which the physicians discovered that the SPRINT FIDELIS Lead, model 6949, had fractured and the shocks were a result of said fracture.

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15.

Between September 2004 and July 2007, Defendants received more than 650 reports of adverse events pertaining to SPRINT FIDELIS Leads filed by physicians and patients. The most frequent complaints were fractured leads caused by the small diameter of the coil and conductor. Fractured leads can deliver unneeded, frightening shocks and potentially life threatening shocks. After analyzing all the leads, defendants concluded that 77 of 125 SPRINT FIDELIS Leads tested (62%) were defective and dangerous due to fracture but defendants continued to market and sell the SPRINT FIDELIS Leads without publishing their findings on testing and analysis.

16.

In a separate study in early 2005, Defendants began a coordinated analysis of more that 25000 patient files who had received SPRINT FIDELIS Leads. Defendants found that their own analysis showed the failure rate of SPRINT FIDELIS Leads was three times higher than the previous ICD lead model but did not publish these findings and continued to manufacturer, market, promote and sell the SPRINT FIDELIS Leads as before.

17.

At all times relevant herein, Defendants knew there was a defect and knew the defect in the SPRINT FIDELIS Lead was attributable to the small diameter of the coil and conductor. Defendants knew that the SPRINT FIDELIS Lead was subject to fracture damage both during and after implant. This information was known to the Defendants prior to implantation in Plaintiff ROSEANN MITCHELL in 2005.

18.

At all times relevant to this action, Defendants knew, that the SPRINT FIDELIS Leads were not safe for the patients for whom they were prescribed and implanted, because the leads fractured and otherwise malfunctioned, and therefore failed to operate in a safe HERSHANDHERSH A Professional Corporation

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and continuous manner, causing injuries from fracture which include, but are not limited to, damages to the blood vessel following insertion, formation of blood clots around the lead itself, additional surgery to explant the fractured leads followed by a second immediate implant surgery, and scarring on fragile compromised heart muscles.

19.

Defendants' representations regarding the performance of the SPRINT FIDELIS Leads, including but not limited to, of the consistency of the performance of the SPRINT FIDELIS Leads, their safety and reliability, were untrue as set forth in their own testing and analysis. The Defendants failed to disclose to physicians, patients or Plaintiff ROSEANN MITCHELL that its leads were subject to fracture causing the injuries herein described.

20.

At all relevant times herein, Defendants continued to promote the SPRINT FIDELIS Lead as safe and effective even when its own internal patient study were reporting high failure rates from fracture while reaping annual sales of well over \$12 billion.

21.

Plaintiff ROSEANN MITCHELL'S SPRINT FIDELIS Lead was explanted on September 11, 2007. The lead system itself had fractured. Plaintiff ROSEANN MITCHELL was, therefore, forced to have an early explant and implant of a new lead system, scarring her already fragile heart, and forcing her to undergo additional and unnecessary complicated surgery. As a result, Plaintiff ROSEANN MITCHELL has suffered severe physical and emotional injuries as a result of the defective SPRINT FIDELIS Lead system in addition to the injuries and damages sustained as a result of the shocks that preceded the explantation.

22.

It was not until October 15, 2007, that the SPRINT FIDELIS Leads model number

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6949, Plaintiff's model, were recalled due to fracture defects leading to severe injury and/or death in patients implanted with this model number.

23.

At all times herein mentioned, the officers and/or directors of the Defendants named herein participated in, authorized and/or directed the production and promotion of the aforementioned products when they knew of the hazards and dangerous propensities of said products, and thereby actively participated in the tortuous conduct that resulted in the injuries suffered by Plaintiffs.

## FIRST CAUSE OF ACTION [Strict Product Liability; Failure to Warn]

24.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-23, inclusive, of this Complaint.

25.

Defendants manufactured, sold and/or distributed SPRINT FIDELIS Leads to Plaintiff ROSEANN MITCHELL to be used in implantation of her ICD.

26.

At all times mentioned herein, SPRINT FIDELIS Leads were and are dangerous and presented a substantial danger to cardiac patients who were implanted with the SPRINT FIDELIS Leads, and these risks and dangers were known or knowable at the time of distribution and implantation in Plaintiff ROSEANN MITCHELL in 2005. Ordinary consumers would not have recognized the potential risks and dangers the SPRINT FIDELIS Leads posed to cardiac patients because its uses was specifically promoted to improve the health of cardiac patients. The SPRINT FIDELIS Leads were used in a way reasonably foreseeable to all Defendants by Plaintiff ROSEANN MITCHELL. Defendants failed to

provide warnings of such risks and dangers to Plaintiff ROSEANN MITCHELL as described herein.

27.

As a result of the fracture of the SPRINT FIDELIS Leads Plaintiff ROSEANN MITCHELL suffered inappropriate shocking leading to the need for dangerous and serious cardiac surgery, required and will continue to require healthcare and services; has incurred and will continue to incur medical and related expenses; has suffered and will continue to suffer mental anguish, diminished capacity for the enjoyment of life, a diminished quality of life, increased risk of premature death, and other such damages.

28.

In doing the acts herein described, the Defendants acted with oppression, fraud and malice, and Plaintiffs are therefore entitled to punitive damages to deter Defendants and others from engaging in similar conduct in the future. Said wrongful conduct was done with advance knowledge, authorization and/or ratification of an officer, director and/or managing agent of the Defendants.

29.

WHEREFORE, Plaintiffs pray for judgment against Defendants as hereinafter set forth.

#### SECOND CAUSE OF ACTION [Strict Liability]

30.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-29, inclusive, of this Complaint.

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The SPRINT FIDELIS Leads were manufactured and/or supplied by the Defendants, and were placed into the stream of commerce by these Defendants in a defective and unreasonably dangerous condition in that the foreseeable risks exceeded the benefits associated with its design of formulation.

32.

Alternatively, the SPRINT FIDELIS Leads manufactured and/or supplied by the Defendants, and each of them, were defective in design or formulation, inadequate warning or instruction and/or inadequate post-marketing warnings or instructions in that when it was placed in the stream of commerce, it was unreasonably dangerous, it was more dangerous than and ordinary consumer would expect and more dangerous than other forms of lead used with ICD's.

33.

As a result of the defective unreasonably dangerous condition of these products manufactured and/or supplied by the Defendants, and each of them, Plaintiff ROSEANN MITCHELL was caused to suffer the herein described injuries and damages.

34.

Defendants acted with conscious and deliberate disregard of the foreseeable harm caused by SPRINT FIDELIS Leads.

35.

Defendants thereby acted with fraud, malice, oppression and a conscious disregard for the Plaintiff and general public's safety, who accordingly requests that the trier of fact, in the exercise of sound discretion, award additional damages for the sake of example and for the purpose of punishing the Defendants for their conduct, in an amount sufficiently large to be an example to others and deter the Defendants and others from engaging in

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similar conduct in the future. The aforesaid wrongful conduct was done with the advance knowledge, authorization, and/or ratification of an officer, director, and/or managing agent of Defendants.

36.

WHEREFORE, Plaintiffs pray for judgment against Defendants as hereinafter set forth.

## THIRD CAUSE OF ACTION [Negligence]

37.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-36, inclusive, of this Complaint.

38.

Defendants, and each of them, and their representatives were manufacturers and/or distributors of SPRINT FIDELIS Leads model 6949. At all times herein, Defendants had a duty to properly manufacture, compound, test, inspect, package, label, distribute, market, examine, maintain supply, provide proper warnings and prepare for use and sell the aforesaid product.

39.

Defendants, and each of them, so negligently and carelessly manufactured, compounded, tested, failed to test, inspected, failed to inspect, packaged, labeled, distributed, recommended, displayed, sold, examined, failed to examine and supplied aforesaid product, that it was dangerous and unsafe for the use and purpose for which it was intended, that is, regulating incorrect heart rates in Plaintiffs and others similarly situated. As a result of the carelessness and negligence of Defendants, Plaintiff ROSEANN MITCHELL had the SPRINT FIDELIS Leads implanted in the manner intended by the

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manufacturer, and, as a result, Plaintiff suffered the injuries and damages described herein.

40.

WHEREFORE, Plaintiffs pray for judgment against Defendants as hereinafter set forth.

## Breach of Implied Warranty

41.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-40, inclusive, of this Complaint.

Defendants impliedly warranted that its SPRINT FIDELIS Leads, which Defendants designed, manufactured, assembled, promoted and sold to Plaintiffs were merchantable and fit and safe for ordinary use. Defendants further impliedly warranted that its SPRINT FIDELIS Leads were fit for the particular purpose of regulating heart rate and rhythm.

43.

Defendants' SPRINT FIDELIS Leads were defective, unmerchantable, and unfit for ordinary use when sold, and unfit for the particular purpose for which they were sold, and subjected Plaintiffs to severe and permanent injuries. Therefore, Defendants breached the implied warranties of merchantability and fitness for a particular purpose when its leads were sold to Plaintiffs, in that the leads are defective and have fractured and otherwise failed to function as represented and intended. 

44.

As a result of Defendants breach of the implied warranties of merchantability and fitness for a particular purpose, Plaintiff ROSEANN MITCHELL has sustained and will continue to sustain the injuries and damages described herein and is therefore entitled to

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compensatory damages.

45.

After Plaintiffs were made aware their injuries were a result of the aforesaid product, SPRINT FIDELIS Leads, Defendants had ample and sufficient notice of the breach of said warranty.

46.

WHEREFORE, Plaintiffs prays for judgment as hereinafter set forth.

## FIFTH CAUSE OF ACTION [Breach of Express Warranty]

47.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-46, inclusive, of this Complaint.

48.

Defendants expressly warranted to Plaintiffs and/or their authorized agents or sales representatives, in publications, and other communications intended for medical patients, and the general public, that the defective leads were safe, effective, fit and proper for their intended use.

49.

Plaintiff ROSEANN MITCHELL, and Plaintiff's physicians reasonably relied upon the skill and judgment of Defendants, and upon said express warranty, in using the aforesaid product. The warranty and representations were untrue in that the product caused severe injury to Plaintiff ROSEANN MITCHELL and was unsafe and, therefore, unsuited for the use in which it was intended and caused Plaintiff ROSEANN MITCHELL to sustain damages and injuries herein alleged.

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50.

As soon as the true nature of the product, and the fact that the warranty and representations were false, were ascertained, said Defendants had ample and sufficient notice of the breach of said warranty.

51.

WHEREFORE, Plaintiffs pray for judgment as hereinafter set forth.

## SIXTH CAUSE OF ACTION [Fraud]

52.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-51, inclusive, of this Complaint.

53:

Defendants falsely and fraudulently represented to Plaintiff ROSEANN MITCHELL, her physicians, and to members of the general public that the aforesaid product was safe, effective, reliable, consistent, and better than the other similar products due to its small diameter for use with implantable ICDs when used in the manner intended by the manufacturer. The representations by said Defendants were in fact, false. The true facts include, but are not limited to the fact that the aforesaid product was not safe to be used in implantation of ICD devices, and were, in fact, dangerous to the health and body of Plaintiff ROSEANN MITCHELL.

54.

When the Defendants made these representations, they knew that they were false. Defendants made said representations with the intent to defraud and deceive plaintiff ROSEANN MITCHELL, with the intent to induce plaintiff to act in the manner herein alleged, that is to use the aforementioned product for implantation with the ICD.

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55.

At the time Defendants made the aforesaid representations Plaintiff ROSEANN MITCHELL took the actions herein alleged, Plaintiff and her physicians were ignorant of the falsity of these representations and reasonably believed them to be true. In reliance upon said representations, plaintiff was induced to, and did, use the aforesaid product as herein described. If Plaintiff ROSEANN MITCHELL had known the actual facts, she would not have taken such action. The reliance of Plaintiff and her physicians upon Defendants representations were justified because said representations were made by individuals and entities that appeared to be in a position to know the true facts.

56.

As a result of Defendants fraud and deceit, Plaintiff was caused to sustain the herein described injuries and damages.

57.

In doing the acts herein alleged, the Defendants acted with oppression, fraud, and malice, and Plaintiff is therefore entitled to punitive damages to deter Defendants and others from engaging in similar conduct in the future. Said wrongful conduct was done with advance knowledge, authorization and/or ratification of an officer, director and/or managing agent of Defendants.

58.

WHEREFORE, Plaintiffs pray for judgment as hereinafter set forth.

#### SEVENTH CLAIM FOR RELIEF [Fraud by Concealment]

59.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-58, inclusive, of this Complaint.

HERSHANDHERSH A Professional Corporation 60.

At all times mentioned herein, Defendants had the duty and obligation to disclose to Plaintiffs and to their physicians, the true facts concerning the aforesaid product, SPRINT FIDELIS Leads, that is, that said product was dangerous and defective, lacking efficacy for its purported use and lacking safety in normal use, and how likely it was to cause serious consequences to users including injuries and possible death. Defendants made the affirmative representations as set forth above to Plaintiffs and their physicians and the general public prior to the date SPRINT FIDELIS Leads were implanted in Plaintiff ROSEANN MITCHELL, while concealing material facts.

61.

At all times herein mentioned, Defendants willfully, and maliciously concealed facts as set forth above from Plaintiffs and their physicians, and therefore, Plaintiffs, with the intent to defraud as herein alleged.

62.

At all times herein mentioned, neither Plaintiffs nor their physicians were aware of the facts set forth above, and had they been aware of said facts, they would not have acted as they did, that is, would not reasonably relied upon said representations of safety and efficacy and utilized the SPRINT FIDELIS Leads for correction of heart rate. Defendants' representations were a substantial factor in Plaintiff utilizing the SPRINT FIDELIS Lead for correction of heart rate and rhythm.

63.

As a result of the concealment of the facts set forth above, Plaintiffs sustained injuries as hereinafter set forth.

64.

In doing the action herein alleged, Defendants acted with oppression, fraud, and malice and Plaintiffs are therefore entitled to punitive damages in an amount reasonably related to Plaintiffs' actual damages, and to Defendant's wealth, and sufficiently large to be an example to others, and to deter these Defendants and others from engaging in similar conduct in the future.

65.

WHEREFORE, Plaintiffs pray for judgment as hereinafter set forth.

## EIGHTH CAUSE OF ACTION [Deceptive Trade Practice]

66.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-65, inclusive, of this Complaint.

67.

Defendants acted deceptive and were misleading in concealing material defects with SPRINT FIDELIS Leads and adverse data from physicians and consumers in order to deceive doctors and patients like Plaintiffs into believing there were significantly fewer adverse events than there really were and are.

68.

Defendants manufactured and promoted both the safety and efficacy of SPRINT FIDELIS Leads to the public and to Plaintiffs' physician for the treatment of regulating heart rate without adverse cardiac events, including but not limited to, terrifying inappropriate defibrillation shocks, failure to deliver appropriate (life-giving) defibrillation therapy and death.

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69.

WHEREFORE, Plaintiffs pray for judgment as hereinafter set forth.

## NINTH CAUSE OF ACTION [Negligent Misrepresentation]

70.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-69, inclusive, of this Complaint.

71.

At all relevant times herein, Defendants represented to Plaintiff ROSEANN MITCHELL and her physicians that the SPRINT FIDELIS Leads were safe to use to correct and regulate heart rate and rhythm knowing that the SPRINT FIDELIS Leads were defective and fractured causing injuries described herein.

72.

The Defendants made the aforesaid representations with no reasonable ground for believing them to be true when defendants own data showed the SPRINT FIDELIS Leads to be defective and dangerous when used in the intended manner.

73.

The aforesaid representations were made to the physician prescribing SPRINT FIDELIS Leads prior to the date it was prescribed to Plaintiffs and their physicians with the intent that Plaintiff and her physicians rely upon such misrepresentations about the safety and efficacy of SPRINT FIDELIS Leads. Plaintiff and her physicians did reasonably rely upon such representations that the aforesaid product was safe for use to aid in the treatment of irregular heart rate.

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74.

The representations by said Defendants to Plaintiff's were false, and thereby caused Plaintiff's injuries described herein.

75.

WHEREFORE, Plaintiffs pray for judgment as hereinafter set forth.

## TENTH CAUSE OF ACTION [Intentional Infliction of Emotional Distress]

76.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-75, inclusive, of this Complaint.

77.

Defendants engaged in extreme and outrageous conduct, knowingly and/or recklessly marketing defective leads, knowingly and/or recklessly concealing a known and potentially fatal defect from Plaintiffs, and knowingly and/or recklessly misrepresenting the quality and usefulness of the SPRINT FIDELIS Leads.

**78.** 

Defendants intended to cause Plaintiffs severe emotional distress, and/or acted with reckless disregard for the Plaintiffs emotional state. As a result of this outrageous conduct and/or reckless disregard for Plaintiff's emotional state, Plaintiffs did, in fact, incur, and continues to incur, severe emotional distress as a result of Defendants' misconduct.

79.

WHEREFORE, Plaintiffs pray for judgment as hereinafter set forth.

# ELEVENTH CLAIM FOR RELIEF ON BEHALF OF URIELA MITCHELL, PATRICIA HINES AND THOMAS MITCHELL [Negligent Infliction of Emotional Distress]

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80.

Plaintiffs hereby incorporate by reference, as if fully set forth herein, each and every allegation contained in Paragraphs 1-79, inclusive, of this Complaint.

81.

Defendants carelessly and negligently manufactured, marketed and sold defective SPRINT FIDELIS Leads to Plaintiff ROSEANN MITCHELL causing her to suffer the herein described injuries from the defective and fractured SPRINT FIDELIS Leads.

82.

Plaintiffs were directly involved in and directly impacted by Defendants carelessness and negligence, in that Plaintiffs were witnesses to multiple episodes of terrifying shocks to Plaintiff ROSEANN MITCHELL.

83.

Defendants' misconduct as alleged above has caused Plaintiffs to suffer severe emotional trauma, physical consequences and long continued emotional disturbance. Plaintiffs are therefore entitled to compensatory damages and equitable and declaratory relief according to proof.

84.

WHEREFORE, Plaintiffs pray for judgment as hereinafter set forth.

#### PRAYER FOR RELIEF

85.

Plaintiffs pray that a judgment be entered in favor of Plaintiffs in such aggregate sum as will fairly and reasonably compensate Plaintiffs for damages arising out of Defendants conduct as described herein. The conduct of Defendants, as alleged herein, was a direct, proximate and producing cause of the damages to Plaintiffs and the following general and specific damages:

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2.	For general damages for Plaintiffs Uriela Mitchell, Thomas Mitchell and
Patricia Hines	s, according to proof;

- 3. For medical, hospital, and incidental expenses, according to proof;
- 4. For loss of earnings and for loss of earning capacity, according to proof;
- 5. For punitive or exemplary damages;
- 6. For such other relief as the Court deems just and proper.

DATED: November 5, 2007

HERSH & HERSH A Professional Corporation

CYNTHIA BROWN Attorneys for Plaintiffs

#### **CT** CORPORATION

A WoltersKluwer Company

Service of Process Transmittal

11/15/2007

CT Log Number 512795580

TO:

Vicki Tersteeg

Medtronic, Inc.
MS: LC300, 710 Medtronic Parkway Minneapolis, MN 55432-5604

RE:

**Process Served in California** 

FOR:

Medtronic, Inc. (Domestic State: MN)

RECEIVED

NOV 1 9 2007

LAW DEPARTMENT

MEDTRO TO THE ABOVE COMPANY AS FOLLOWS:

TITLE OF ACTION:

Roseann Mitchell, et al., Pltfs. vs. Medtronic, Inc., et al., Dfts.

DOCUMENT(S) SERVED:

Summons, Complaint, Cover Sheet, Notice to Plaintiff, Statement Form(s),

Attachment(s), Stipulation Form

**COURT/AGENCY:** 

San Francisco County- San Francisco, Superior Court, CA

Case # CGC07468894

NATURE OF ACTION:

Product Liability Litigation - Personal Injury - Sprint Fidelis Leads

ON WHOM PROCESS WAS SERVED:

C T Corporation System, Los Angeles, CA

DATE AND HOUR OF SERVICE:

By Process Server on 11/15/2007 at 14:20

APPEARANCE OR ANSWER DUE:

Within 30 days after service - file written response // 4/4/2008 at 9:00 a.m. - Case

Management Conference

ATTORNEY(S) / SENDER(S):

Nancy Hersh Hersh & Hersh 208 Opera Plaza 601 Van Ness Avenue San Francisco, CA 94102 415 441 5544

**ACTION ITEMS:** 

Telephone, Vicki Tersteeg , 763-505-2671 SOP Papers with Transmittal, via Fed Ex Priority Overnight , 798310819350 Email Notification, Vicki Tersteeg VICKI.ANN.TERSTEEG@MEDTRONIC.COM Email Notification, Anne Sederstrom ANNE.SEDERSTROM@MEDTRONIC.COM

SIGNED: PER ADDRESS: C T Corporation System

Nancy Flores 818 West Seventh Street Los Angeles, CA 90017 213-337-4615

TELEPHONE:

Page 1 of 1/JC

Information displayed on this transmittal is for CT Corporation's record keeping purposes only and is provided to the recipient for quick reference. This information does not constitute a legal opinion as to the nature of action, the amount of damages, the answer date, or any information contained in the documents themselves. Recipient is responsible for interpreting said documents and for taking appropriate action. Signatures on certified mail receipts confirm receipt of package only, not contents.

Daputy Clark

County Superior Court NOV 0 52007 GORDON PARK-LI, Clerk

CASE MANAGEMENT CONFERENCE SET

APR 0 4 2008 - 9MAM

DEPARTMENT 212

OSEANN MITCHELL, URIELA	) CASE NUMBER
ITCHELL, THOMAS MITCHELL	)
nd PATRICIA HINES,	) COMPLAINT FOR DAMAGES AND
	) DEMAND FOR JURY TRIAL
Plaintiffs,	)
· • • • • • • • • • • • • • • • • • • •	) [PRODUCTS LIABILITY]
5.	)
	) 1. Strict Liability-Failure to Warn
MEDTRONIC, INC., McKESSON	) 2. Strict Product Liability
ORPORATION and DOES ONE	) 3. Negligence
brough FIFTY, inclusive,	) 4. Breach of Implied Warranty
	) 5. Breach of Express Warranty
Defendants.	) 6. Fraud
	7. Fraud by Concealment
	) 8. Deceptive Trade Practice
	9. Negligent Misrepresentation
•	) 10. Intentional Infliction of Emotional
	) Distress
•	) 11. Negligent Infliction of Emotional
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	) Distress

COMPLAINT FOR DAMAGES AND DEMAND FOR JURY TRIAL

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PERIOR COURT OF CALIFORNIA, COUNTY OF SA	n Francisco .	
STREET ADDRESS: 400 McAllister St	reet	GORDON PARKLI, Clerk
MAILING ADDRESS:	04107	DEBORAH STEPPE
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CASE NAME: ROSEANN MITCHELL, BT A	L. v. MEDIRONIC, INC., ET AL.	
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	Drugs (38)	Other complaint (not specified above) (42)
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Unless this is a collections case under it	ule 3.740 or a complex case, this cover s	sheet will be used for statistical purposes only.
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Judicial Countil of Galilomia Control (Rev. July 1, 2007) MagFooths (509) 595-4382

CASE NUMBER: CGC-07-468894 ROSEANN MITCHELL et al. VS. MEDTRONIC, INC. et al

### **NOTICE TO PLAINTIFF**

A Case Management Conference is set for

DATE:

APR-04-2008

TIME:

9:00AM

PLACE:

Department 212

400 McAllister Street

San Francisco, CA 94102-3680

All parties must appear and comply with Local Rule 3.

CRC 3.725 requires the filing and service of a case management statement form CM-110 no later than 15 days before the case management conference.

However, it would facilitate the issuance of a case management order without an appearance at the case management conference if the case management statement is filed, served and lodged in Department 212 twenty-five (25) days before the case management

Plaintiff must serve a copy of this notice upon each party to this action with the summons and complaint. Proof of service subsequently filed with this court shall so state.

#### **ALTERNATIVE DISPUTE RESOLUTION POLICY REQUIREMENTS**

IT IS THE POLICY OF THE SUPERIOR COURT THAT EVERY CIVIL CASE PARTICIPATE IN EITHER MEDIATION, JUDICIAL OR NON-JUDICIAL ARBITRATION, THE EARLY SETTLEMENT PROGRAM OR SOME SUITABLE FORM OF ALTERNATIVE DISPUTE RESOLUTION PRIOR TO A MANDATORY SETTLEMENT CONFERENCE OR TRIAL. (SEE LOCAL RULE 4)

Plaintiff must serve a copy of the Alternative Dispute Resolution Information Package on each defendant along with the complaint. All counsel must discuss ADR with clients and opposing counsel and provide clients with a copy of the Alternative Dispute Resolution Information Package prior to filing the Case Management Statement.

[DEFENDANTS: Attending the Case Management Conference does not take the place of filing a written response to the complaint. You must file a written response with the court within the time limit required by law. See Summons.]

Superior Court Alternative Dispute Resolution Coordinator 400 McAllister Street, Room 103 San Francisco, CA 94102 (415) 551-3876

See Local Rules 3.6, 6.0 C and 10 D re stipulation to commissioners acting as temporary judges

	CM-110
ATTORNEY OR PARTY WITHOUT ATTORNEY (Name. Since But number, and iddress).	FOR COURT USE OPLY
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1. Party or parties (answer one):	•
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b. This statement is submitted jointly by parties (names):	
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b. The cross-complaint, if any, was filed on (date):	•
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3. Service (to be answered by plaintiffs and cross-complainants only)	•
a. All parties named in the complaint and cross-complaint have been served,	or have appeared, or have been dismissed,
b. The following parties named in the complaint or cross-complaint	••
(1) have not been served (specify names and explain why not):	•
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c. The following additional parties may be added (specify names, nature of in	volvement in case, and the date by which
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C.	Dates on which parties or attorneys will not	be available for trial (spe	cify dates and ex	plain reasons for un	svailability);
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9.	Party represented:				
	Additional representation is described in A	Attachment 8.		•	
). Pred	erance   This case is antitled to preference (specify	r code section):			
0. Altı 9.	ernative Dispute Resolution (ADR)  Counsel has has hes not pro reviewed ADR options with the client.	ovided the ADR Informati	ion package ident	ilied in rule 3,221 to	Une client and has
b.	All parties have agreed to a form of Al	DR. ADR will be complete	ed by (date):		
C,	The case has gone to an ADR process		•		
M. FIR IR	V. January 1, 2007		•		Page 7 of 4

PLAINTIFF/PETITIONER;	CASE HUNDER	•
DEFENDANT/RESPONDENT:		
O. d. The party or parties are willing to participate in (check all that apply):  (1) Mediation  (2) Monbinding judicial arbitration under Code of Civil Procedure section 1 arbitration under Cal. Rules of Court, rule 3,822)	141.12 (discovery to close	15 days before
(3) Nonbinding judicial arbitration under Code of Civil Procedure section 1	141.12 (discovery to rema	in open until 30 day
before trial; order required under Cal. Rules of Court, rule 3.822)  (4) Binding judicial arbitration		•
(5) Binding private arbitration		
(6) Neutral case evaluation		•
(7) L Other (specify):	•	•
e. This matter is subject to mandatory judicial arbitration because the amount	in controversy does not ex	ceed the statutory i
<ol> <li>Plaintiff elects to refer this case to judicial arbitration and agrees to limit reco Procedure section 1141.11.</li> </ol>	overy to the amount specifi	ed in Code of Civil
g. This case is exempt from judicial arbitration under rule 3.811 of the Californi	is Rules of Court (specific a	zemolihe),
		······································
. Settlement conference		•
The party or parties are willing to participate in an early settlement conference (s	specify when):	
	to and seconds	•
. Insurance		
a. Insurance camer, if any, for party fling this statement (name):		•
b Reservation of rights: Yes No		÷
c. Coverage issues will significantly affect resolution of this case (explain):	•	
and the second of the second factors of the second factors in		
	•	
Junediction	•	•
Indicate any matters that may affect the court's jurisdiction or processing of this case, a	and describe the status.	
Bankrupicy Other (specify):		
'Status;		•
Related cases, consolidation, and coordination  a There are companion, underlying or related cases		* * * * * * * * * * * * * * * * * * * *
a. There are companion, underlying, or related cases.  (1) Name of case:		•
(2) Name of court		Ç.
(3) Case number:	•	
(4) Status:  Additional cases are described in Attachment 14a,	4	
	f	
b. A motion to consolidate coordinate will be filed by (	name party):	•
Bilurcation		* * * * * * * * * * * * * * * * * * * *
The party or parties intend to file a motion for an order bifurcating, severing, or co action (specify moving party, type of motion, and reasons):	eel pniwollof ent pnibalion	ues or causes of
Other mations		
The party or parties expect to file the following motions before trial (specify maying	a nadu hmm et eestes	/ m
	a haith' tàhe o <u>t wotton</u> ' auc	issues);
·		
	•	

	•			•	CM-110
	PLAINTIFF/PETITIONER:			CASE NUMBER:	
ĎE	FENDANTIRESPONDENT:		٠.		
17,		ve completed all discovery. I will be completed by the date spe	icified (describe all ani	: icipated discovery);	•
ė	Party	Description		<u>Date</u>	
	•	٠.		•	
	c The following discovery	issues are anticipated (spacify):		•	
18.	of Civil Procedure section	se (i.e., the amount demanded is ions 80 through 98 will apply to this se and a motion to withdraw the coff checked, explain specifically who case):	s case. asa from the economic	: Illigation procedures or for ac	iditional
19,	Other leaves The party or parties reques conference (specify):	st that the following additional mat	lars be considered or c	letermined at the case manag	enent
20,		ve met and conferred with all parti ):	es on all subjects requ	ired by rule 3,724 of the Califo	rnia Ruies
	<ul> <li>b. After meeting and conferring (specify):</li> </ul>	as required by rule 3.724 of the C	California Rules of Cou	rt, the parties agree on the foll	owing
21.	Case management orders Previous case management ord	ers in this case are (check one):	none a	ttached as Attachment 21.	•
22.	Total number of pages attached	(if any):			
rais COI	n completely familiar with this cas sed by this statement, and will pos ference, including the written auti te:	ises the authority to enter into stip	pulations and assur pulations on these issu	wary and ADR, as well as othe ses at the time of the case man	er issues nagement
			•		
	(TYPÉ OR PRINT N	WE)	jsk.	NATURE OF PARTY OR ATTORNEY)	· · · · · · · · · · · · · · · · · · ·
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~	440 77				Page 4 of 4



## Superior Court of California County of San Francisco

## Judicial Mediation Program

Introducing a new court alternative dispute resolution program that provides judicial mediation of complex civil cases

The Judicial Mediation program offers mediation of complex civil litigation by a San Francisco Superior Court judge familiar with the area of the law that is the subject of the controversy. Cases that will be considered for participation in the program include, but are not limited to professional malpractice, construction, employment, insurance coverage disputes, mass torts and complex commercial litigation. Judicial mediation offers civil litigants the opportunity to engage in early mediation of a case shortly after filing the complaint in an effort to resolve the matter before substantial funds are expended. This program may also be utilized at anytime throughout the litigation process. The panel of judges currently participating in the program includes:

The Honorable David L. Ballati The Honorable Anne Bouliane The Honorable Ellen Chaitin The Honorable John J. Conway The Honorable Robert L. Dondero The Honorable Emest H. Goldsmith The Honorable Curtis B. A. Karnow The Honorable Patrick J. Mahoney

The Honorable Tomar Mason The Honorable James J. McBride The Honorable Kevin M. McCarthy The Honorable John E. Munter The Honorable Ronald Evans Quidachay The Honorable A. James Robertson, II The Honorable Mary E. Wiss

Parties interested in judicial mediation should file the Stipulation to Alternative Dispute Resolution form attached to this packet indicating a joint request for inclusion in the program and deliver a courtesy copy to Dept. 212. A preference for a specific judge may be indicated. The court Alternative Dispute Resolution Coordinator will facilitate assignment of cases that qualify for the program.

Note: Space is limited. Submission of a stipulation to judicial mediation does not guarantee inclusion in the program. You will receive written notification from the court as to the outcome of your application.

> Superior Court Alternative Dispute Resolution 400 McAllister Street, Room 103, San Francisco, CA 94102 (415) 551-3876

Page 42 of 85

## Alternative Dispute Resolution (ADR) Information Package

Document 1-2

## Alternatives to Trial

# Here are some other ways to resolve a civil dispute.

The plaintiff must serve a copy of the ADR information package on each defendant along with the complaint. (CRC 201.9(c))

> Superior Court of California County of San Francisco

#### Introduction

Did you know that most civil lawsuits settle without a trial?

And did you know that there are a number of ways to resolve civil disputes without having to sue somebody?

Document 1-2

These alternatives to a lawsuit are known as alternative dispute resolutions (ADR). The most common forms of ADR are mediation, arbitration and case evaluation. There are a number of other kinds of ADR as well.

In ADR, trained, impartial persons decide disputes or help parties decide disputes themselves. These persons are called neutrals. For example, in mediation, the neutral is the mediator. Neutrals normally are chosen by the disputing parties or by the court. Neutrals can help parties resolve disputes without having to go to court.

ADR is not new. ADR is available in many communities through dispute resolution programs and private neutrals.

#### Advantages of ADR

ADR can have a number of advantages over a lawsuit.

- ADR can be speedier. A dispute often can be resolved in a matter of months. even weeks, through ADR, while a lawsuit can take years.
- ADR can save money. Court costs, attorneys fees, and expert fees can be saved.
- ADR can permit more participation. The parties may have more chances to tell their side of the story than in court and may have more control over the outcome.
- ADR can be flexible. The parties can choose the ADR process that is best for them. For example, in mediation the parties may decide how to resolve their dispute.
- ADR can be cooperative. This means that the parties having a dispute may work together with the neutral to resolve the dispute and agree to a remedy that makes sense to them, rather than work against each other.

ADR-1 1/06 (bc) Page 2 ADR can reduce stress. There are fewer, if any, court appearances. And because ADR can be speedier, and save money, and because the parties are normally cooperative, ADR is easier on the nerves. The parties don't have a lawsuit hanging over their heads for years.

Document 1-2

ADR can be more satisfying. For all the above reasons, many people have reported a high degree of satisfaction with ADR.

Because of these advantages, many parties choose ADR to resolve a dispute, instead of filing a lawsuit. Even when a lawsuit has been filed, the court can refer the dispute to a neutral before the parties' position harden and the lawsuit becomes costly. ADR has been used to resolve disputes even after a trial, when the result is appealed.

## Disadvantages of ADR

ADR may not be suitable for every dispute.

- If ADR is binding, the parties normally give up most court protections, including a decision by a judge or jury under formal rules of evidence and procedure, and review for legal error by an appellate court.
- There generally is less opportunity to find out about the other side's case with ADR than with litigation. ADR may not be effective if it takes place before the parties have sufficient information to resolve the dispute.
- The neutral may charge a fee for his or her services.
- If a dispute is not resolved through ADR, the parties may have to put time and money into both ADR and a lawsuit.
- Lawsuits must be brought within specified periods of time, known as statutes of limitation. Parties must be careful not to let a statute of limitations run out while a dispute is in an ADR process.

## **ALTERNATIVE DISPUTE RESOLUTION PROGRAMS** Of the San Francisco Superior Court

Document 1-2

"It is the policy of the Superior Court that every noncriminal, nonjuvenile case participate either in an early settlement conference, mediation, arbitration, early neutral evaluation or some other alternative dispute resolution process prior to a mandatory settlement conference or trial." (Superior Court Local Rule 4)

This guide is designed to assist attorneys, their clients and self-represented litigants in complying with San Francisco Superior Court's alternative dispute resolution ("ADR") policy. Attorneys are encouraged to share this guide with clients. By making informed choices about dispute resolution alternatives, attorneys, their clients and self-represented litigants may achieve a more satisfying resolution of civil disputes.

The San Francisco Superior Court currently offers three ADR programs for civil matters; each program is described below:

- Judicial arbitration 1).
- Mediation 2)
- The Early Settlement Program (ESP) in conjunction with the San Francisco Bar Association.

#### **JUDICIAL ARBITRATION**

## Description

In arbitration, a neutral "arbitrator" presides at a hearing where the parties present evidence through exhibits and testimony. The arbitrator applies the law to the facts of the case and makes an award based upon the ments of the case. When the Court orders a case to arbitration it is called judicial arbitration. The goal of arbitration is to provide parties with an adjudication that is earlier, faster, less formal, and usually less expensive than a trial. Upon stipulation of all parties, other civil matters may be submitted to judicial arbitration.

Although not currently a part of the Court's ADR program, civil disputes may also be resolved through private arbitration. Here, the parties

ADR-1 1/06 (bc) Page 4

voluntarily consent to arbitration. If all parties agree, private arbitration may be binding and the parties give up the right to judicial review of the arbitrator's decision. In private arbitration, the parties select a private arbitrator and are responsible for paying the arbitrator's fees.

Document 1-2

#### Operation

Pursuant to CCP 1141.11 and Local Rule 4, all civil actions in which the amount in controversy is \$50,000 or less, and no party seeks equitable relief, shall be ordered to arbitration. A case is ordered to arbitration after the Case Management Conference. An arbitrator is chosen from the Court's Arbitration Panel. Most cases ordered to arbitration are also ordered to a pre-arbitration settlement conference. Arbitrations are generally held between 7 and 9 months after a complaint has been filed. Judicial arbitration is not binding unless all parties agree to be bound by the arbitrator's decision. Any party may request a court trial within 30 days after the arbitrator's award has been filed.

#### Cost

There is no cost to the parties for judicial arbitration or for the prearbitration settlement conference.

#### **MEDIATION**

## Description

Mediation is a voluntary, flexible, and confidential process in which a neutral third party "mediator" facilitates negotiations. The goal of mediation is to reach a mutually satisfactory agreement that resolves all or part of the dispute after exploring the significant interests, needs, and priorities of the parties in light of relevant evidence and the law.

Although there are different styles and approaches to mediation, most mediations begin with presentations of each side's view of the case. The mediator's role is to assist the parties in communicating with each other, expressing their interests, understanding the interests of opposing parties, recognizing areas of agreement and generating options for resolution. Through questions, the mediator aids each party in assessing the strengths and weaknesses of their position.

Page 5 ADR-1 1/06 (bc)

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A mediator does not propose a judgment or provide an evaluation of the merits and value of the case. Many attorneys and litigants find that mediation's emphasis on cooperative dispute resolution produces more satisfactory and enduring resolutions. Mediation's non-adversarial approach is particularly effective in disputes in which the parties have a continuing relationship, where there are multiple parties, where equitable relief is sought, or where strong personal feelings exist.

Document 1-2

#### Operation

San Francisco Superior Court Local Court Rule 4 provides three different voluntary mediation programs for civil disputes. An appropriate program is available for all civil cases, regardless of the type of action or type of relief sought.

To help litigants and attorneys identify qualified mediators, the Superior Court maintains a list of mediation providers whose training and experience have been reviewed and approved by the Court. The list of court approved mediation providers can be found at www.sfgov.org/courts. Litigants are not limited to mediators on the court list and may select any mediator agreed upon by all parties. A mediation provider need not be an attorney.

Local Rule 4.2 D allows for mediation in lieu of judicial arbitration, so long as the parties file a stipulation to mediate within 240 days from the date the complaint is filed. If settlement is not reached through mediation, a case proceeds to trial as scheduled.

#### **Private Mediation**

The Private Mediation program accommodates cases that wish to participate in private mediation to fulfill the court's alternative dispute resolution requirement. The parties select a mediator, panel of mediators or mediation program of their choice to conduct the mediation. The cost of mediation is borne by the parties equally unless the parties agree otherwise.

Parties in civil cases that have not been ordered to arbitration may consent to private mediation at any point before trial. Parties willing to submit a matter to private mediation should indicate this preference on the Stipulation to Alternative Dispute Resolution form or the Case Management Statement (CM-110). Both forms are attached to this packet.

ADR-1 1/06 (bc) Page 6

## Mediation Services of the Bar Association of San Francisco

The Mediation Services is a coordinated effort of the San Francisco Superior Court and The Bar Association of San Francisco (BASF) in which a court approved mediator provides three hours of mediation at no charge to the parties. It is designed to afford civil litigants the opportunity to engage in early mediation of a case shortly after filing the complaint, in an effort to resolve the matter before substantial funds are expended on the litigation process. Although the goal of the program is to provide the service at the outset of the litigation, the program may be utilized at anytime throughout the litigation process.

Document 1-2

The mediators participating in the program have been pre-approved by the court pursuant to strict educational and experience requirements.

After the filing of the signed Stipulation to Alternative Dispute Resolution form included in this ADR package the parties will be contacted by BASF. Upon payment of the \$200 per party administration fee, parties select a specific mediator from the list of court approved mediation providers. The hourly mediator fee beyond the first three hours will vary depending on the mediator selected. Waiver of the administrative fee based on financial hardship is available.

A copy of the Mediation Services rules can be found on the BASF website . at www.sfbar.org, or you may call BASF at 415-782-8913

#### Judicial Mediation

The Judicial Mediation program is designed to provide early mediation of complex cases by volunteer judges of the San Francisco Superior Court. Cases considered for the program include construction defect, employment discrimination, professional malpractice, insurance coverage, toxic torts and industrial accidents.

Parties interested in judicial mediation should file the Stipulation to Alternative Dispute Resolution form attached to this packet indicating a joint request for inclusion in the program. A preference for a specific judge may be indicated. The court Alternative Dispute Resolution Coordinator will coordinate assignment of cases that qualify for the program.

ADR-1 1/06 (bc) Page 7

#### Cost

Generally, the cost of Private Mediation ranges from \$200 per hour to \$400 per hour and is shared equally by the parties. Many mediators are willing to adjust their fees depending upon the income and resources of the parties. Any party who meets certain eligibility requirements may ask the court to appoint a mediator to serve at no cost to the parties.

The Mediation Services of the Bar Association of San Francisco provides three hours of mediation time at no cost with a \$200 per party administrative fee.

There is no charge for participation in the Judicial Mediation program.

#### **EARLY SETTLEMENT PROGRAM**

#### Description

The Bar Association of San Francisco, in cooperation with the Court, offers an Early Settlement Program ("ESP") as part of the Court's settlement conference calendar. The goal of early settlement is to provide participants an opportunity to reach a mutually acceptable settlement that resolves all or part of the dispute. The two-member volunteer attorney panel reflects a balance between plaintiff and defense attorneys with at least 10 years of trial experience.

As in mediation, there is no set format for the settlement conference. A conference typically begins with a brief meeting with all parties and counsel, in which each is given an opportunity to make an initial statement. The panelists then assist the parties in understanding and candidly discussing the strengths and weaknesses of the case. The Early Settlement Conference is considered a "quasi-judicial" proceeding and, therefore, is not entitled to the statutory confidentiality protections afforded to mediation.

#### Operation

Civil cases enter the ESP either voluntarily or through assignment by the Court. Parties who wish to choose the early settlement process should indicate this preference on the status and setting conference statement.

ADR-I 1/06 (bc) Page 8

If a matter is assigned to the ESP by the Court, parties may consult the ESP program materials accompanying the "Notice of the Early Settlement Conference" for information regarding removal from the program.

Document 1-2

Participants are notified of their ESP conference date approximately 4 months prior to trial. The settlement conference is typically held 2 to 3 months prior to the trial date. The Bar Association's ESP Coordinator informs the participants of names of the panel members and location of the settlement conference approximately 2 weeks prior to the conference date.

Local Rule 4.3 sets out the requirements of the ESP. All parties to a case assigned to the ESP are required to submit a settlement conference statement prior to the conference. All parties, attorneys who will try the case, and insurance representatives with settlement authority are required to attend the settlement conference. If settlement is not reached through the conference, the case proceeds to trial as scheduled.

#### Cost

All parties must submit a \$200 generally non-refundable administrative fee to the Bar Association of San Francisco. Parties who meet certain eligibility requirements may request a fee waiver. For more information, please contact the ESP Coordinator at (415) 982-1600.

For further information about San Francisco Superior Court ADR programs or dispute resolution alternatives, please contact:

> Superior Court Alternative Dispute Resolution Coordinator, 400 McAllister Street, Room 103 San Francisco, CA 94102 (415) 551-3876

or visit the Superior Court Website at http://sfgov.org/site/courts\_page.asp?id=3672

ADR-1 1/06 (bc)

## SUPERIOR COURT OF CALIFORNIA

•	•		
	<b>v.</b>	Plaintiff	Case No STIPULATION TO ALTERNATIVE DISPUTE RESOLUTION
-	••	Defendant .	
The resolution p	parties hereby : rocess:	Upulate that this action	shall be submitted to the following alternative dispute
ما المالية	BASF Early		Mediation Services of BASF U Judicial Mediation Judge Judge
Plain	ntiff(s) and Defe	ndant(s) further agree a	s follows:
Plain	ntiff(s) and Defe		
		ndant(s) further agree a	
		ndant(s) further agree a	Attorney Executing Stipulation Signature of Party or Attornay
		ndant(s) further agree a	
Name of Party	Stipulating	Name of Party or	Attorney Executing Stipulation Signature of Party or Attornay
Name of Party  J Plaintiff  Name of Party	Stipulating	Name of Party or  Cross-detendant  Name of Party or	Attorney Executing Stipulation Signature of Party or Attorney  Dated:
Name of Party Plaintiff Name of Party	Stipulating  Defendent  Stipulating  Defendant	Name of Party or Cross-defendant  Name of Party or Cross-defendant  Cross-defendant	Attorney Executing Stipulation Signature of Party or Attorney  Dated:
Name of Party	Stipulating  Defendent  Stipulating  Defendant	Name of Party or Cross-defendant  Name of Party or Cross-defendant  Cross-defendant	Attorney Executing Stipulation Signature of Party or Attorney  Dated:

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REED SMITH LLP

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2	REED SMITH LLP
3	1999 Harrison St., Suite 2400 Oakland, CA 94612
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6	REED SMITH LLP
	355 South Grand Avenue, Suite 2900
7	Los Angeles, CA 90071-1514
	Telephone: 213.457.8000
8	Facsimile: 213.457.8080
9	Attorneys for Defendant Medtronic, Inc.

ENDORSED
Superior Count of California
County of San Francisco

DEC 1 3 2007

GORDON PARK-LI, Clerk

BY: MARIA SANCHEZ

Deputy Clerk

## SUPERIOR COURT OF THE STATE OF CALIFORNIA FOR THE COUNTY OF SAN FRANCISCO

ROSEANN MITCHELL, et al.,

Plaintiffs,

vs.

MEDTRONIC, INC., et al.,

Defendants.

No.: CGC-07-468894

DEFENDANT MEDTRONIC, INC.'S ANSWER TO COMPLAINT

Compl. Filed: November 5, 2007

Defendant Medtronic, Inc. ("Medtronic"), hereby answers Plaintiffs' Complaint as follows:

Answering Plaintiffs' unverified Complaint, Medtronic denies each and every allegation contained therein and denies that by reason of any act or omission by Medtronic, its agents, employees, or independent contractors, Plaintiffs were injured or damaged in any sum, or at all.

#### AFFIRMATIVE DEFENSES

Medtronic, while reserving the right to assert other defenses as discovery proceeds, and without assuming the burden of proof when the burden of proof rests on Plaintiff, asserts the following separate and independent affirmative defenses in further opposition to the Complaint:

#### FIRST AFFIRMATIVE DEFENSE

Plaintiffs' Complaint, and each count and claim contained therein, fails to state a claim upon which relief can be granted.

#### SECOND AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, by the applicable statutes of limitation and/or the doctrine of laches.

#### THIRD AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, by the doctrines of estoppel, unclean hands and waiver.

#### FOURTH AFFIRMATIVE DEFENSE

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Plaintiffs' claims are barred, in whole or in part, by the doctrines of res judicata, collateral estoppel, issue preclusion and/or claim preclusion.

#### FIFTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because the Food and Drug Administration has exclusive or primary jurisdiction over the matters asserted in the Complaint.

#### SIXTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are preempted, in whole or in part, by federal laws and regulations, including without limitation those governing the labeling, advertisement and sale of medical devices.

#### SEVENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are preempted, in whole or in part, by the deference state law gives to discretionary actions by the Food and Drug Administration under the Food, Drug and Cosmetic Act, and the Medical Device Amendments thereto.

#### **EIGHTH AFFIRMATIVE DEFENSE**

Plaintiffs' claims are barred, in whole or in part, because there is no private right of action under the Food, Drug and Cosmetic Act for claims such as those asserted by Plaintiff.

#### NINTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because at all times relevant to the Complaint, Medtronic met or exceeded the requisite standard of care.

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#### TENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because Medtronic did not owe any duty to Plaintiffs.

#### **ELEVENTH AFFIRMATIVE DEFENSE**

Plaintiffs' claims are barred, in whole or in part, because no act or omission on Medtronic's part caused or contributed to the alleged injuries and damages described in the Complaint.

#### TWELFTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because Plaintiffs' alleged injuries, if any, were the result of intervening and/or superseding causes.

#### THIRTEENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, by Plaintiffs' comparative or contributory fault or negligence.

#### FOURTEENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because no action or inaction by Medtronic was the proximate cause of Plaintiffs' alleged damages, if any.

#### FIFTEENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because Plaintiffs' alleged damages, if any,

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were caused in whole or in part by the acts or omissions of Plaintiffs or third parties over whom Medtronic had no control or authority.

#### SIXTEENTH AFFIRMATIVE DEFENSE

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Plaintiffs' claims are barred, in whole or in part, because Plaintiff assumed the risk of her alleged injuries, if any, and engaged in the activities alleged in the Complaint after giving her informed consent.

#### SEVENTEENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, by the learned intermediary doctrine, because Medtronic provided adequate warnings to learned intermediaries.

#### EIGHTEENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because any alleged injuries or damages sustained by Plaintiff may have been caused by the misuse or abuse of Medtronic's products by Plaintiff or other persons.

#### NINETEENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because Medtronic's products may have been substantially changed after they left Medtronic's control and before Plaintiffs suffered any alleged injuries or damages.

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Plaintiffs' claims are barred, in whole or in part, because any alleged injuries or damages sustained by Plaintiff may have been caused by the alteration and/or method of implantation and/or maintenance of Medtronic's products after they left Medtronic's control.

#### TWENTY-FIRST AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because any alleged injuries or damages sustained by Plaintiff may have been the direct result of Plaintiffs' pre-existing medical conditions, sub-medical conditions, natural causes, or the result of other circumstances over which Medtronic had no control and for which Medtronic is not responsible.

#### TWENTY-SECOND AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because any alleged injuries or damages sustained by Plaintiffs may be the result of an unforeseeable series of events over which Medtronic had no control, and as such, constitutes acts of God for which Medtronic cannot be held liable.

#### TWENTY-THIRD AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because any alleged injuries or damages sustained by Plaintiffs may be the result of idiosyncratic reactions by Plaintiffs that were not reasonably foreseeable and for which Medtronic cannot be held liable.

#### TWENTY-FOURTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because the risks associated with the use of

the medical devices at issue, if any, are outweighed by the utility and benefits such devices provide.

#### TWENTY-FIFTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because the medical device at issue was not unreasonably dangerous or defective, was suitable for the purpose for which it was intended, and was distributed with adequate and sufficient warnings.

#### TWENTY-SIXTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because the methods, standards and techniques of designing, manufacturing and labeling the medical devices at issue complied with and conformed to the generally recognized state of the art at the time such devices were designed, manufactured and labeled.

#### TWENTY-SEVENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because Plaintiffs fail to assert a feasible safer design for Medtronic's products alleged to be defective.

#### TWENTY-EIGHTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because Medtronic acted reasonably and integrated good faith at all times material herein, based on all relevant facts and circumstances known by Medtronic at the time it acted.

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#### THIRTIETH AFFIRMATIVE DEFENSE

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Plaintiffs' claims are barred, in whole or in part, because there was no defect in the device at issue at the time it left Medtronic's possession.

#### THIRTY-FIRST AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because to the extent Plaintiffs allege that Medtronic failed to warn about alleged defects in the device at issue or other alleged causes of Plaintiffs' injuries or damages, if any, the doctors and other health care providers associated with the device were or should have been aware of any risks or hazards associated with it, and to the extent that such doctors and health care providers failed to advise, inform or warn Plaintiff of such risks or hazards, Medtronic cannot be held responsible.

#### THIRTY-SECOND AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, to the extent that they have failed to plead actual injury.

#### THIRTY-THIRD AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, to the extent that the alleged injuries are too remote from Medtronic's conduct to state a claim.

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#### THIRTY-FOURTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, by the economic loss doctrine.

#### THIRTY-FIFTH AFFIRMATIVE DEFENSE

Medtronic adopts and incorporates by reference as if fully set out herein any and all defenses which are or may become available to it under the Restatement (Second) of Torts § 402A and all comments thereto, and the Restatement (Third) of Torts §§ 1-21 and comments thereto.

#### THIRTY-SIXTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, by the principles set forth in the Restatement (Second) of Torts § 388, Comment n, and any similar doctrines and/or principle in the Restatement (Third) of Torts.

#### THIRTY-SEVENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, by the principles set forth in the Restatement (Second) of Torts § 402A, Comment k, and the Restatement (Third) of Torts: Products Liability § 6.

#### THIRTY-EIGHTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because to the extent Plaintiffs alleges failure to warn by Medtronic, the doctors and other health care providers who were associated with the device at issue, or were or should have been aware of any risk and/or hazard that Plaintiff alleges rendered the device defective and allegedly caused Plaintiffs' damages, if any, failed to warn Plaintiff of such risks and/or hazards.

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#### THIRTY-NINTH AFFIRMATIVE DEFENSE

Plaintiffs' warranty-based claims are barred, in whole or in part, because Medtronic did not make or breach any warranties that are applicable to Plaintiff.

#### FORTIETH AFFIRMATIVE DEFENSE

Plaintiffs' warranty-based claims are barred, in whole or in part, by Plaintiffs' failure to give proper or timely notice of any alleged defect or breach of warranty.

#### FORTY-FIRST AFFIRMATIVE DEFENSE

Plaintiffs' warranty-based claims are barred, in whole or in part, because Plaintiffs were not in privity with Medtronic.

#### FORTY-SECOND AFFIRMATIVE DEFENSE

Plaintiffs' warranty-based claims are barred, in whole or in part, by any and all express conditions, disclaimers, modifications or exclusions made by Medtronic.

#### FORTY-THIRD AFFIRMATIVE DEFENSE

Plaintiffs' warranty-based claims are barred, in whole or in part, by Plaintiffs' lack of reliance on any such warranties.

#### FORTY-FOURTH AFFIRMATIVE DEFENSE

21.

Plaintiffs' warranty-based claims are barred, in whole or in part, by Plaintiffs' failure to

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satisfy all conditions precedent or subsequent to the enforcement of any such alleged warranties.

#### FORTY-FIFTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because the promotion of products sold or manufactured by Medtronic is protected by the First Amendment of the United States Constitution and similar provisions in applicable state constitutions.

#### FORTY-SIXTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, to the extent that they lack standing to pursue the claims alleged against Medtronic.

#### FORTY-SEVENTH AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, in whole or in part, because Medtronic did not violate any statute or law, as alleged by Plaintiff.

#### FORTY-EIGHTH AFFIRMATIVE DEFENSE

Because have failed to plead fraud claims with particularity, their fraud claims must be dismissed.

#### FORTY-NINTH AFFIRMATIVE DEFENSE

Plaintiffs' claims under state consumer fraud and unlawful or deceptive trade practice acts are barred to the extent that Plaintiffs have not alleged any misrepresentation or misleading statement with the specificity required.

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Plaintiffs' claims are barred, in whole or in part, because Medtronic is entitled to the benefit of all defenses and presumptions contained in, or arising from, any rule of law or statute of any state whose substantive law controls the action.

#### FIFTY-FIRST AFFIRMATIVE DEFENSE

Plaintiffs' claims are barred, reduced and/or limited pursuant to applicable statutory and common law regarding limitation of awards, caps on recovery and setoffs.

#### FIFTY-SECOND AFFIRMATIVE DEFENSE

Any verdict or judgment that might be recovered by Plaintiffs must be reduced by those amounts that have already indemnified Plaintiffs, or will in the future with reasonable certainty indemnify Plaintiffs, in whole or in part, for any past or future claimed economic loss, from any collateral source including but not limited to insurance, social security, workers' compensation or employee benefit programs.

#### FIFTY-THIRD AFFIRMATIVE DEFENSE

In the event that Plaintiffs have sustained damages as alleged in the Complaint, which Medtronic denies, discovery or investigation may reveal that Plaintiffs' claims are barred or reduced to the extent Plaintiffs failed to mitigate any damages allegedly sustained.

#### FIFTY-FOURTH AFFIRMATIVE DEFENSE

Plaintiffs' claims for punitive or exemplary damages are barred or reduced by applicable law

or statute, or in the alternative, are unconstitutional insofar as they violate the due process protections afforded by the United States Constitution, the excessive fines clause of the Eighth Amendment to the United States Constitution, and any other applicable provisions of the United States Constitution or any applicable state constitution. Any law, statute or other authority purporting to permit the recovery of punitive or exemplary damages in this case is unconstitutional, facially and/or as applied to Medtronic.

FIFTY-FIFTH AFFIRMATIVE DEFENSE

Plaintiffs' claims for punitive or exemplary damages are barred, in whole or in part, because such damages are not recoverable for the causes of action set forth in the Complaint, or in the alternative, the allegations of each cause of action in the Complaint are legally insufficient to support a claim for punitive or exemplary damages.

#### FIFTY-SIXTH AFFIRMATIVE DEFENSE

Plaintiffs' claims for punitive or exemplary damages are barred, in whole or in part, because Medtronic did not act with the requisite level of conduct to be subjected to or that would otherwise support any punitive or exemplary damages in this action.

#### FIFTY-SEVENTH AFFIRMATIVE DEFENSE

Plaintiffs are not entitled to attorneys' fees under any act or theory forming the basis of any of Plaintiffs' claims.

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one or more other claims asserted in the Complaint or (2) do not satisfy the prerequisites for such

claims as bystanders.

FIFTY-NINTH AFFIRMATIVE DEFENSE

Medtronic hereby gives notice that it intends to rely upon any other defenses that may become available or apparent during the discovery proceedings in this matter and hereby reserves its right to amend its Answer to assert any such defense.

WHEREFORE, having fully answered and defended, Medtronic d prays for judgment as follows:

- 1. That Plaintiffs take nothing by the Complaint;
- 2. That judgment be entered for Medtronic and against Plaintiffs on each and every claim set forth in the Complaint;
  - 3. That Medtronic recover its costs of suit; and
  - 4. For such other and further relief as the Court deems just and proper.

DATED: December 13, 2007

REED SMITH LLP

Sonja S. Weissman

Dana A. Blanton Attorneys for Defendant

Medtronic, Inc.

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1	PROOF OF SERVICE				
2	I am a resident of the State of California, over the age of eighteen years, and not a				
3	party to the within action. My business address is REED SMITH LLP,				
4	1999 Harrison Street, Suite 2400, Oakland, CA 94612-3572. On December 13, 2007, I served the				
5	following document(s) by the method indicated below:				
6	DEFENDANT MEDTRONIC, INC.'S ANSWER TO COMPLAINT				
7	by transmitting via facsimile on this date from fax number +1 510 273 8832 the document(s)				
8	listed above to the fax number(s) set forth below. The transmission was completed before 5:00 PM and was reported complete and without error. The transmission report, which is attached to this proof of service, was properly issued by the transmitting fax machine.				
10	Service by fax was made by agreement of the parties, confirmed in writing. The transmitting fax machine complies with Cal.R.Ct 2003(3).				
11	by placing the document(s) listed above in a sealed envelope with postage thereon fully prepaid, in the United States mail at Oakland, California addressed as set forth below. I am				
12	readily familiar with the firm's practice of collection and processing of correspondence for mailing. Under that practice, it would be deposited with the U.S. Postal Service on that same				
13	day with postage thereon fully prepaid in the ordinary course of business. I am aware that on motion of the party served, service is presumed invalid if the postal cancellation date or				
14	postage meter date is more than one day after the date of deposit for mailing in this Declaration.				
15	by placing the document(s) listed above in a sealed envelope(s) and by causing personal				
16	delivery of the envelope(s) to the person(s) at the address(es) set forth below. A signed proof of service by the process server or delivery service will be filed shortly.				
17 18	by personally delivering the document(s) listed above to the person(s) at the address(es) set forth below.				
19	by placing the document(s) listed above in a sealed envelope(s) and consigning it to an				
20	express mail service for guaranteed delivery on the next business day following the date of consignment to the address(es) set forth below. A copy of the consignment slip is attached to this proof of service.				
21	by transmitting via email to the parties at the email addresses listed below:				
22					
23	Attorneys for Plaintiff				
24	Nancy Hersh, Esq.				
	Mark E. Burton, Jr., Esq.  Rachel Abrams, Esq.				
25	Cynthia Brown, Esq.				
26	Hersh & Hersh				
Ó	2080 Opera Plaza				
27	601 Van Ness Avenue				
28	San Francisco, CA 94102 Telephone: (415) 441-5544				

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REED SMITH LLP

I declare under penalty of perjury under the laws of the State of California that the above is true and correct. Executed on December 13, 2007, at Oakland, California.

Trisha Suzette Hooper

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UNITED STATES DISTRICT COURT WESTERN DISTRICT OF WASHINGTON AT SEATTLE

IN RE: PHENYLPROPANOLAMINE (PPA) PRODUCTS LIABILITY LITIGATION,

MDL NO. 1407

ORDER DENYING PLAINTIFF'S MOTION TO REMAND

This document relates to:

Barnett, et al. v. American Home Products Corp., et al., No. C02-423R

THIS MATTER comes before the court on the motion of plaintiffs to remand the case to state court in Mississippi. reviewed the papers filed in support of and in opposition to this motion, the court rules as follows:

#### I. BACKGROUND

Plaintiffs purchased a variety of over-the-counter drugs including, but not limited to, products sold under the trade names "Robitussin," "Alka-Seltzer Plus," "Dimetapp," "Tavist D," "BC," "Triaminic," "Contac," "Comtrex," and "Equate Tussin CF." All of these products contained the ingredient phenylpropanolamine ("PPA"). The individuals later consumed the medication and suffered unidentified types of injuries. In June 2001, plaintiffs filed an amended complaint in Mississippi state court linking the PPA in the medicine with the injuries sustained.

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manufacturers and distributors of PPA-containing products, as well as several retail stores that sold those products. One of the stores named as a defendant, Bill's Dollar Stores, Inc., d/b/a Bill's Dollar Store ("Bill's Dollar Store"), is a Mississippi corporation. Two of the six total plaintiffs purchased PPA-containing products from Bill's Dollar Store.

The complaint alleges numerous causes of action against both

Defendants removed the complaint to federal court alleging that plaintiffs fraudulently joined Bill's Dollar Store. Plaintiffs moved to remand to state court. The case was later transferred to this court as part of a multi-district litigation ("MDL").

#### II. ANALYSIS

A plaintiff cannot defeat federal jurisdiction by fraudulently joining a non-diverse party. As an MDL court sitting in the Ninth Circuit, this court applies the Ninth Circuit's fraudulent joinder standard to the motion to remand. See, e.g., In reduction Drugs Prods. Liab. Litiq., 220 F. Supp. 2d 414, 423 (E.D. Pa. 2002); In reduction Firestone, 204 F. Supp. 2d 1149, 1152 n.2 (S.D. Ind. 2002); In reduction for Tobacco/Gov'tal Health Care Costs Litiq., 100 F. Supp. 2d 31, 34 n.1 (D. D.C. 2000); In reduction for the supp. 2d 31, 34 n.1 (D. D.C. 2000); In reduct

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Defendants assert the misjoinder of these plaintiffs' claims and request that the court sever and deny remand as to the four plaintiffs who did not purchase any products from Bill's Dollar Store, or from any other Mississippi store. However, because, as discussed below, the court denies remand as to all plaintiffs named in this action, the court need not address the question of misjoinder at this time.

Ford Motor Co. Bronco II Prods. Liab. Litig., MDL-991, 1996 U.S. Dist. LEXIS 6769, at \*2-4 (E.D. La. May 16, 1996). Under this standard, joinder of a non-diverse party is deemed fraudulent "'[i]f the plaintiff fails to state a cause of action against a resident defendant, and the failure is obvious according to the settled rules of the state.' Morris v. Princess Cruises. Inc., 236 F.3d 1061, 1067 (9th Cir. 2001) (quoting McCabe v. General Foods Corp., 811 F.2d 1336, 1339 (9th Cir. 1987)).

The propriety of removal to federal court is determined from the allegations in the complaint at the time of removal. See Ritchev v. Upjohn Drug Co., 139 F.3d 1313, 1318 (9th Cir. 1998) However, in the case of fraudulent joinder, the defendant "'is entitled to present the facts showing the joinder to be fraudulent." Id. (quoting McCabe, 811 F.2d at 1339). See also Morris

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<sup>&</sup>lt;sup>2</sup>See generally Menowitz v. Brown, 991 F.2d 36, 40-41 (2d Cir. 1993); In Re Korean Airlines Disaster, 829 F.2d 1171, 1174-76 (D.C. Cir. 1987).

<sup>&#</sup>x27;However, as a practical matter, application of the Fifth Circuit's fraudulent joinder standard would not alter the court's conclusion. See Badon v. RJR Nabisco, Inc., 224 F.3d 382, 393 (5th Cir. 2000) (remand is denied where there is "no reasonable basis for predicting that plaintiffs might establish liability . . . against the in-state defendants.") For example, recent MDL courts utilized fraudulent joinder standards similar, and in one case identical, to the Fifth Circuit's standard in deeming Mississippi pharmacies and their employees fraudulently joined for reasons similar to those expressed in this opinion. re Diet Drugs Prods, Liab, Litig., 220 F. Supp. 2d at 423-24 (noting that there had been "a pattern of pharmacies being named in complaints, but never pursued to judgment, typically being voluntarily dismissed at some point after the defendants' ability to remove the case has expired"); In re Rezulin Prods. Liab. Litig., 133 F. Supp. 2d 272, 279 & n.3, 288-92 (S.D.N.Y. 2001).

236 F.3d at 1067-68 (citing <u>Cavallini v. State Farm Mut. Auto.</u>

<u>Ins. Co.</u>, 44 F.3d 256, 263 (5th Cir. 1995) for the proposition
that the court may "'pierc[e] the pleadings'" and consider

"summary judgment-type evidence.")

Defendants allege that plaintiffs fraudulently joined Bill's Dollar Store, while plaintiffs claim the existence of legitimate causes of action against Bill's Dollar Store, including products liability, negligence, misrepresentation, and implied warranty claims. The parties also argue as to the relevance of a bank-ruptcy petition filed by Bill's Dollar Store prior to the filing of this suit.

## A. <u>Products Liability</u>

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The complaint contains failure to warn and design defect allegations pursuant to the Mississippi Products Liability Act.

Miss. Code Ann. § 11-1-63. Under the Products Liability Act, plaintiff must show that at the time the product left the control of the manufacturer or seller, it was defective in failing to contain adequate warnings or instructions, and/or was designed in a defective manner. Miss. Code Ann. § 11-1-63 (a) (i) (2)-(3). Plaintiff must also show that the manufacturers and sellers knew, or in light of reasonably available knowledge or the exercise of reasonable care should have known, about the danger that caused the alleged damage. Miss. Code Ann. § 11-1-63 (c) (i), (f) (i).

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<sup>&</sup>lt;sup>4</sup>See also <u>Huff v. Shopsmith. Inc.</u>, 786 So.2d 383, 387 (Miss. 2001) ("With the adoption of 11-1-63, common law strict liability, as laid out in <u>State Stove Mfg. Co. v. Hodges</u>, 189 So.2d 113

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First, the complaint utilizes the plural "defendants" in a number of allegations that one could not reasonably interpret to include Bill's Dollar Store. See, e.g., Louis v. Wyeth-Ayerst Pharm.. Inc., No. 5:00CV102LN, slip op. at 5-9 (S.D. Miss. Sep. 25, 2000) (finding products liability allegations lodged against "defendants" conclusory where there was no factual support for conclusion that Mississippi pharmacies had knowledge or reason to know of alleged dangers associated with various diet drugs).

(Miss. 1966), is no longer the authority on the necessary elements of a products liability action.")

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See also In re Diet Drugs Prods. Liab. Litig., 220 F. Supp. 2d at 424 (finding complaints, including failure to warn, negligence, breach of warranty, and strict liability claims, devoid of specific allegations against Mississippi pharmacies and "filled instead with general statements levied against all defendants, which most properly can be read as stating claims against drug manufacturers."); In re Rezulin Products Liab. Litig., 133 F. Supp. 2d at 291 (finding improper joinder in case where Mississippi pharmacies were lumped in with manufacturers and acts alleged, including failure to warn, breach of warranty, and fraud, were attributed to "'defendants' generally", but never connected to the pharmacies); accord Badon, 224 F.3d at 391-93 ("While the amended complaint does often use the word

For example, the complaint describes "defendants" as members of the Non-Prescription Drug Manufacturers Association ("NDMA"). Through this association, "defendants" purportedly participated in numerous discussions relating to the safety of PPA over the past two decades, had representatives sit on the NDMA PPA Task Force, and funded relevant studies. In other words, plaintiffs, in significant part, demonstrate "defendants" knowledge as to risks allegedly posed by PPA through activities engaged in by manufacturer defendants alone.

Indeed, while "defendants" are alleged to have been aware or to have had responsibility for awareness of numerous scientific journal articles, incident reports, medical textbooks, and other reports containing information as to risks of PPA consumption, general medical practitioners are excluded from this awareness and described as being not "fully informed." The complaint supplies no factual support for a conclusion that a dollar store possessed medical and scientific knowledge beyond that possessed by medical practitioners.

Second, the complaint specifically lays the responsibility for allegedly concealing dangers posed by PPA on the manufacturer defendants. For example, the complaint alleges that the manufacturer defendants concealed material facts regarding PPA through product packaging, labeling, advertising, promotional campaigns

<sup>&#</sup>x27;defendants,' frequently it is evident that such usage could not be referring to the 'Tobacco Wholesalers.'"; finding conspiracy allegations against Louisiana defendants entirely general).

and materials, and other methods. This allegation directly undermines and contradicts the idea that Bill's Dollar Store had knowledge or reason to know of alleged defects. Seg, e.g., Louis, slip op. at 4-5 (finding complaint's "major theme" to consist of the "manufacturers' intentional concealment of the true risks of the drug(s), coupled with dissemination through various media of false and misleading information of the safety of the drug(s) at issue, [which belied] any suggestion of knowledge, or reason to know by [the] resident defendants.") Cf. In respectively. In the Rezulin Products Liab, Littic., 133 F. Supp. 2d 272, 290 (S.D.N.Y. 2001) (finding Mississippi pharmacies facing failure to warn claims fraudulently joined where "the theory underlying the complaints [was] that the manufacturer defendants hid the dangers of Rezulin from plaintiffs, the public, physicians, distributors and pharmacists — indeed from everyone.")

In sum, the court concludes that one could not reasonably read the complaint to support the idea that the retailer defendants had knowledge or reason to know of any dangers allegedly associated with PPA. Indeed, reading the complaint as a whole, this allegation reveals itself as directed towards the manufacturer defendants alone. As such, the court finds that plaintiffs fail to state a products liability cause of action against Bill's Dollar Store.

The complaint once alludes to an "alternative" breach of express warranty claim under the Products Liability Act. See Miss. Code Ann. \$ 11-1-63 (a) (i) (4) (requiring a showing that the

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# B. Negligence and Misrepresentation

The complaint alleges negligence and misrepresentation by Bill's Dollar Store. A negligence cause of action also requires a showing of knowledge or reason to know on the part of the seller. See, e.g., R. Clinton Constr. Co. v. Bryant & Reaves. Inc., 442 F. Supp. 838, 851 (N.D. Miss. 1977) ("The rule is well settled that in order to fasten liability upon a party for negligence, it must be shown by a preponderance of the evidence that he knew or through the exercise of reasonable care should have known that his selection of a [product] would cause damage to his customer.")" A misrepresentation cause of action requires

seller breached an express warranty or failed to conform to other express factual representations upon which the claimant relied). However, the products liability allegations go on to touch solely upon failure to warn and design defect claims. Because the complaint lacks any factual basis for support of a breach of express warranty claim against Bill's Dollar Store, the court also finds this bare allegation insufficient to support remand.

7 Accord Louis, slip op. at 3-4 & n.3 ("[K] nowledge, or a reason to know, is also a necessary requisite for any claim of failure to warn or negligence that a plaintiff might undertake to assert extraneous to a claim under the Products Liability Act itself (assuming solely for the sake of argument that such a claim could exist)."); Cadillac Corp. v. Moore, 320 So.2d 361, 365 (Miss. 1975) (discussing negligence in "vendor/purchaser" context and stating that "fault on the part of a defendant so as to render him liable is to be found in action or nonaction, accompanied by knowledge, actual or implied, of the probable result of his conduct.") Cf. Moore v. Memorial Hosp. of Gulfport, 825 So.2d 658, 664-66 (Miss. 2002) (extending "learned intermediary" doctrine to pharmacists in case involving prescription drug, and holding no actionable negligence claim could exist against a pharmacy unless a plaintiff indisputably informed the pharmacy of health problems which contraindicated the use of the drug in question, or the pharmacist filled

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a plaintiff to show:

(1) a representation; (2) its falsity; (3) its materiality; (4) the speaker's knowledge of its falsity or ignorance of its truth; (5) the speaker's intent that the representation should be acted upon by the hearer and in the manner reasonably contemplated; (6) the hearer's ignorance of its falsity; (7) the hearer's reliance on its truth; (8) the hearer's right to rely thereon; and (9) the hearer's consequent and proximate injury.

Johnson v. Parke-Davis, 114 F. Supp. 2d 522, 525 (S.D. Miss. 2000) (citing Allen v. Mac Tools, Inc., 671 So.2d 636, 642 (Miss. 1996)).

Again, the court finds that the general and contradictory allegations in the complaint do not support the existence of any knowledge or reason to know on the part of Bill's Dollar Store to support a negligence cause of action. The court finds the complaint similarly bereft of any factual support for the idea that Bill's Dollar Store made any misrepresentations whatsoever to plaintiffs regarding the PPA-containing products. See, e.g., Johnson, 114 F. Supp. 2d at 525 ("Suffice it to say that Plaintiffs have no proof . . . that any of the named [Mississippi] representatives made any representations directly to any of the Plaintiffs. Thus, none of the Plaintiffs was the 'hearer' of any of the sales representatives' alleged misrepresentations."; finding plaintiffs had no cause of action for misrepresentation). Instead, as discussed above, the complaint attributes this

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prescriptions in quantities inconsistent with the recommended dosage guidelines).

behavior to the manufacturing defendants alone. As such, the court also finds that plaintiffs fail to state negligence and misrepresentation causes of action against Bill's Dollar Store.

## C. <u>Implied Warranty</u>

The complaint also alleges that Bill's Dollar Store breached implied warranties of merchantability and fitness for particular purpose. See Miss. Code Ann. \$\$ 75-2-314, 315. The complaint accuses "defendants" of breaching the implied warranty of merchantability in failing to adequately label containers and packages containing PPA, and because the products sold failed to conform to promises or affirmations of facts made on the containers or labels. See Miss. Code Ann. \$ 75-2-314 (2)(e)-(f). The complaint accuses both manufacturers and sellers of breaching the implied warranty of fitness for particular purpose where they had reason to know of the particular use of the products, and the purchasers relied on the sellers' skill or judgment in selecting and furnishing suitable and safe products. See Miss. Code Ann. \$ 75-2-315.

In order to recover for breach of implied warranty, a buyer "must within a reasonable time after he discovers or should have discovered any breach notify the seller of breach or be barred from any remedy." Miss. Code Ann. § 75-2-607 (3) (a); accord C.R. Daniels. Inc. v. Yazoo Mfg. Co., 641 F. Supp. 205, 210-11 (S.D. Miss. 1986); Gast v. Rogers-Dingus Chevrolet, 585 So. 2d 725, 730-31 (Miss. 1991). Here, the complaint contains no indication that plaintiffs provided Bill's Dollar Store with any notice as ORDER Page - 10 -

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to an alleged breach of warranty prior to the institution of this lawsuit.

Additionally, with respect to the merchantability claim, the complaint contains no factual support for a conclusion that Bill's Dollar Store was in any way involved with the labeling and/or packaging of the products at issue. Instead, the complaint alleges that the manufacturer defendants concealed material facts regarding PPA through product packaging and labeling.

The court likewise finds plaintiffs' fitness for particular purpose allegation insufficient. "Mississippi does not recognize an implied warranty of fitness for a particular purpose when the good is purchased for the ordinary purpose of a good of that kind." Farris v. Coleman Co., 121 F. Supp. 2d 1014, 1018 (N.D. Miss. 2000) (fitness for particular purpose claim failed where plaintiff purchased cooler to keep food and beverages cold - the ordinary purpose for which a cooler is used). Here, plaintiffs attested that they purchased PPA-containing products to remedy their "cold, flu, sinus and/or allergy symptoms" - the ordinary purpose of these medications.

Therefore, for the reasons stated above, the court finds that plaintiffs fail to state implied warranty causes of action against Bill's Dollar Store.

#### D. Bankruptcy

Bill's Dollar Store filed a bankruptcy petition in February 2001, several months prior to the filing of plaintiffs' complaint. The filing of the bankruptcy petition operates as a stay ORDER

on judicial or other proceedings brought against Bill's Dollar store that were or could have commenced prior to the commencement of the bankruptcy proceeding. <u>See</u> 11 U.S.C. **S** 362(a): <u>In re</u> <u>Cajun Elec. Power Co-Op. Inc.</u>, 185 F.3d 446, 457 (5th Cir. 1999).

Plaintiffs argue that the automatic stay poses no barrier to relief given that they were unaware of the bankruptcy petition at the time they filed their complaint, and because they anticipate that the Bankruptcy Court will agree to their pending request to lift the stay. However, whether or not plaintiffs knew of the petition and whether or not the stay may later be lifted, the fact remains that, at the time plaintiffs filed their complaint, the stay operated to prohibit their lawsuit. As noted above, the court determines jurisdiction based on the claims as stated at the time of removal. As such, the court finds the existence of the stay at the time of filing serves as an additional reason to deny remand of this matter to state court. Cf. Ritchey, 139 F.3d at 1319-20 (denying remand where the statute of limitations had expired at the time plaintiff filed the complaint).

### III. CONCLUSION

The court concludes that plaintiffs fail to state a cause of action against the only non-diverse defendant, and that the

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<sup>\*</sup>Unlike in a number of other cases transferred to this MDL, the defendants here did not supply the court with any summary judgment-type evidence to establish the retailer defendant's fraudulent joinder. However, the court nonetheless finds that a plain reading of the complaint does not allow a conclusion that plaintiffs state a cause of action against Bill's Dollar Store.

failure is obvious according to the settled rules of Mississippi. As such, the court finds Bill's Dollar Store fraudulently joined and DENIES plaintiff's motion to remand the case to the state courts of Mississippi.

DATED at Seattle, Washington this 26th day of November, 2002.

BARBARA JACOBS ROTHSTEIN UNITED STATES DISTRICT JUDGE

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Filed 12/14/2007 Page 84 of 85

Case 3:07-cv-06352-MMC

Affadvit Of